NRC FORM 374 U.S. NUCLEAR REGU	LATORY COMMISSION PAGE <u>1</u> OF <u>4</u> PAGES Amendment No. 21			
MATERIALS LICENSE Pursuant to the Atomic Energy Act of 1954, as amended, the Energy Reorganization Act of 1974 (Public Law 93-438), and Title 10, Code of Federal Regulations, Chapter I, Parts 30, 31, 32, 33, 34, 35, 36, 39, 40, and 70, and in reliance on statements and representations heretofore made by the licensee, a license is hereby issued authorizing the licensee to receive, acquire, possess, and transfer byproduct, source, and special nuclear material designated below; to use such material for the purpose(s) and at the place(s) designated below; to deliver or transfer such material to persons authorized to receive it in accordance with the regulations of the applicable Part(s). This license shall be deemed to contain the conditions specified in Section 183 of the Atomic Energy Act of 1954, as amended, and is subject to all applicable rules, regulations, and orders of the Nuclear Regulatory Commission now or hereafter in effect and to any conditions specified below.				
Licensee	In accordance with letter dated			
1. Memorial Hospital of Sweetwater County	March 19, 2013 3. License number 49-17940-01 is amended in its entirety to read as follows:			
2. 1200 College Drive	4. Expiration date June 30, 2014			
2. 1200 College Drive Rock Springs, Wyoming 82901	5. Docket No. 030-13672 Reference No.			
 Byproduct, source, and/or special 7. Chemical and/o nuclear material 	or physical form 8. Maximum amount that licensee may possess at any one time under this license			
A. Any byproduct material A. Any permitted by 10 CFR 35.100	A. As needed			
B. Any byproduct material permitted by 10 CFR 35.200	B. As needed			
C. lodine-131 permitted by 10 CFR 35.300	C. 100 millicuries total			
D. Any byproduct material permitted by 10 CFR 31.11	ge Kits D. 10 millicuries total			
Products A3410; N Scientific	purces (IsotopeE.300 millicuries per source and 1 curie totalorth AmericantotalModel MED 3601; or roducts Laboratoriess			

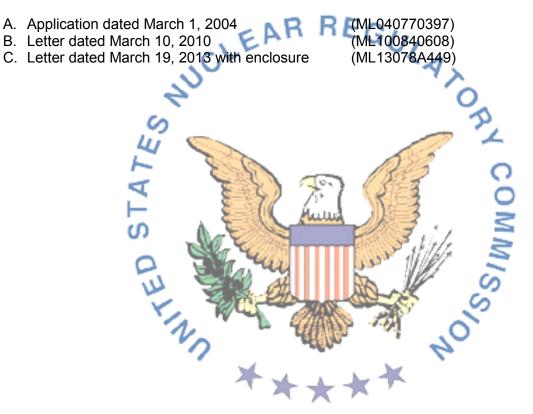
- 9. Authorized use:
 - A. Any uptake, dilution and excretion study permitted by 10 CFR 35.100.
 - B. Any imaging and localization study permitted by 10 CFR 35.200.
 - C. Any sodium iodide lodine-131 use permitted by 10 CFR 35.300 for which the patient can be released under the provisions of 10 CFR 35.75.
 - D. <u>In vitro</u> studies.
 - E. For use in ADAC Laboratories Vantage Non-uniform Attenuation Correction System for instrument calibration and quality control.

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		CONDITIONS	6			
10.		nsed material may be used and/or stored only at the e, Rock Springs, Wyoming.	licensee's facilities located at 1200 College			
11.	The	Radiation Safety Officer for this license is Keith Carr	nahan, RT N.			
12.	Lice	nsed material is only authorized for use by, or under	the supervision of:			
	A. I a	ndividuals permitted to work as an authorized user a ccordance with 10 CFR 35.13 and 35.14.	nd/or authorized nuclear pharmacist in			
	В. Т	he following individuals are authorized users for the	material and medical uses indicated:			
	<u>/</u>	Authorized Users Material and Use	O'S			
	Frederick Matti, M.D. 35.100; 35.200; 31.11; oral administration of sodium iodide I-131 in quantities less than or equal to 33 millicuries; and gadolinium-153 for attenuation correction					
	ſ		gadolinium-153 for attenuation correction			
13.	mate	dition to the possession limits in Item 8, the licensee rial to quantities below the minimum limit specified i rance for decommissioning.				
14.	For s	ealed sources not associated with 10 CFR Part 35 i	use, the following conditions apply:			
	A.	Sealed sources shall be tested for leakage and/or of intervals specified in the certificate of registration is Commission under 10 CFR 32.210 or under equivalent	sued by the U.S. Nuclear Regulatory			
	B.	Notwithstanding Paragraph A of this Condition, sea particles shall be tested for leakage and/or contami				
	C.	In the absence of a certificate from a transferor indi- intervals specified in the certificate of registration is Commission under 10 CFR 32.210 or under equiva- the transfer, a sealed source received from another the test results received.	sued by the U.S. Nuclear Regulatory lent regulations of an Agreement State, prior to			
	D.	Sealed sources need not be tested if they contain or radioactive gas; or the half-life of the isotope is 30 of 100 microcuries of beta- and/or gamma-emitting material.	days or less; or they contain not more than			

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	E. Sealed sources need not be tested if they are in storage and are not being used; however, when they are removed from storage for use or transferred to another person and have not been tested within the required leak test interval, they shall be tested before use or transfer. No sealed source shall be stored for a period of more than 10 years without being tested for leakage and/or contamination.						
	F. The leak test shall be capable of detecting the presence of 0.005 microcurie (185 becquerels) of radioactive material on the test sample. If the test reveals the presence of 0.005 microcurie (185 becquerels) or more of removable contamination, a report shall be filed with the U.S. Nuclear Regulatory Commission in accordance with 10 CFR 30.50(c)(2), and the source shall be removed immediately from service and decontaminated, repaired, or disposed of in accordance with Commission regulations. The report shall be filed within 5 days of the date the leak test result is known with the U.S. Nuclear Regulatory Commission, Region IV, 1600 East Lamar Boulevard, Arlington, Texas 76011-4511, ATTN: Director, Division of Nuclear Materials Safety. The report shall specify the source involved, the test results, and corrective action taken.						
	G. Tests for leakage and/or contamination, including leak test sample collection and analysis, shall be performed by the licensee or by other persons specifically licensed by the U.S. Nuclear Regulatory Commission or an Agreement State to perform such services.						
	H.	Records of leak test results shall be kept in units of	microcuries and shall be maintained for 3 years.				
15.	5. The licensee shall conduct a physical inventory every 6 months, or at other intervals approved by the U.S. Nuclear Regulatory Commission, to account for all sources and/or devices received and possessed under the license. Records of inventories shall be maintained for 3 years from the date of each inventory and shall include the radionuclides, quantities, manufacturer's name and model numbers, and the date of the inventory.						
16.	Sealed sources containing licensed material shall not be opened or sources removed from source holders by the licensee.						
17.		icensee is authorized to transport licensed material FR Part 71, "Packaging and Transportation of Radic					

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18. Except as specifically provided otherwise in this license, the licensee shall conduct its program in accordance with the statements, representations, and procedures contained in the documents, including any enclosures, listed below. This license condition applies only to those procedures that are required to be submitted in accordance with the regulations. Additionally, this license condition does not limit the licensee's ability to make changes to the radiation protection program as provided for in 10 CFR 35.26. The U.S. Nuclear Regulatory Commission's regulations shall govern unless the statements, representations, and procedures in the licensee's application and correspondence are more restrictive than the regulations.



FOR THE U.S. NUCLEAR REGULATORY COMMISSION

Date <u>March 20, 2013</u>

/RA/

By .

Roberto J. Torres, Senior Health Physicist Nuclear Materials Safety Branch B Region IV Arlington, Texas 76011-4511